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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/647,140	09/27/2000	Gary Kruh	FCCC 98-02	5331
110	7590 12/12/2002			
DANN DORFMAN HERRELL & SKILLMAN			EXAMINER	
SUITE 720 1601 MARKET STREET		Ì	CHEN, SHIN LIN	
PHILADELPHIA, PA	HIA, PA 19103-2307	Ì	ART UNIT	PAPER NUMBER
			1632	11
			DATE MAILED: 12/12/2002	lγ

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary

Application No. 09/647,140 Applicant(s)

Kruh et al.

Examiner

Shin-Lin Chen

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	The MAILING DATE of this communication appears	on the cover sheet with the correspondence address		
Period 1	for Reply			
THE	ORTENED STATUTORY PERIOD FOR REPLY IS SET MAILING DATE OF THIS COMMUNICATION.	TO EXPIRE MONTH(S) FROM no event, however, may a reply be timely filed after SIX (6) MONTHS from the		
mailing	date of this communication.			
	period for reply specified above is less than thirty (30) days, a reply within the period for reply is specified above, the maximum statutory period will apply a	e statutory minimum of thirty (30) days will be considered timely. nd will expire SIX (6) MONTHS from the mailing date of this communication.		
	to reply within the set or extended period for reply will, by statute, cause the ply received by the Office later than three months after the mailing date of the	· · ·		
	patent term adjustment. See 37 CFR 1.704(b).			
Status				
1) ∐	Responsive to communication(s) filed on	·		
2a) 🗌	This action is FINAL . 2b) \square This act	on is non-final.		
3) 🗆	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11; 453 O.G. 213.			
Disposi	tion of Claims			
4) 💢	Claim(s) <u>1-59</u>	is/are pending in the application.		
4	a) Of the above, claim(s)	is/are withdrawn from consideration.		
5) 🗌	Claim(s)	is/are allowed.		
_	Claim(s)			
	Claim(s)			
		are subject to restriction and/or election requirement.		
	tion Papers			
9) 🗌	The specification is objected to by the Examiner.			
10)	The drawing(s) filed on is/are	a) accepted or b) objected to by the Examiner.		
	Applicant may not request that any objection to the d	rawing(s) be held in abeyance. See 37 CFR 1.85(a).		
11)	The proposed drawing correction filed on	is: a) \square approved b) \square disapproved by the Examiner.		
	If approved, corrected drawings are required in reply t	o this Office action.		
12)	The oath or declaration is objected to by the Exami	ner.		
Priority	under 35 U.S.C. §§ 119 and 120			
13)	Acknowledgement is made of a claim for foreign pr	iority under 35 U.S.C. § 119(a)-(d) or (f).		
a) [☐ All b)☐ Some* c)☐ None of:			
	1. \square Certified copies of the priority documents hav	e been received.		
	2. \square Certified copies of the priority documents hav	e been received in Application No		
	application from the International Bure			
🖂	ee the attached detailed Office action for a list of the			
14)∟.	Acknowledgement is made of a claim for domestic			
a) ∟ 15\□		• •		
15)	Acknowledgement is made of a claim for domestic	priority under 35 U.S.C. 33 120 and/or 121.		
Attachm	ent(s) tice of References Cited (PTO-892)	4) 🗆 🕒		
	tice of Draftsperson's Patent Drawing Review (PTO-948)	4) Interview Summary (PTO-413) Paper No(s).		
	ormation Disclosure Statement(s) (PTO-1449) Paper No(s).	5) Notice of Informal Patent Application (PTO-152)		
"" لـــا ،"		6) Other:		

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1. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-7, 45-51 and 56-58, drawn to an isolated nucleic acid molecule having the sequence of SEQ ID No. 1 (MOAT-B), a vector comprising said nucleic acid molecule, a host cell containing said nucleic acid molecule, and a method for screening a test compound for inhibition of MOAT mediated transport by using a host cell comprising SEQ ID No. 1.

Group II, claim(s) 8-10, drawn to an antibody specific to MOAT-B.

Group III, claim(s) 11-17, 45-51 and 56-58, drawn to an isolated nucleic acid molecule having the sequence of SEQ ID No. 3 (MOAT-C), a vector comprising said nucleic acid molecule, a host cell containing said nucleic acid molecule, and a method for screening a test compound for inhibition of MOAT mediated transport by using a host cell comprising SEQ ID No. 3.

Group IV, claim(s) 18-20, drawn to an antibody specific to MOAT-C.

Group V, claim(s) 21 and 59, drawn to an oligonucleotide between 10 and 200 nucleotides, which specifically hybridize with a protein translation initiation site in a nucleotide

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sequence encoding SEQ ID No. 4, and a kit containing oligonucleotide primer sequence for amplification.

Group VI, claim(s) 22 and 59, drawn to an oligonucleotide between 10 and 200 nucleotides, which specifically hybridize with a protein translation initiation site in a nucleotide sequence encoding SEQ ID No. 2, and a kit containing oligonucleotide primer sequence for amplification.

Group VII, claim(s) 23-29, 45-51 and 56-58, drawn to an isolated nucleic acid molecule having the sequence of SEQ ID No. 5 (MOAT-D), a vector comprising said nucleic acid molecule, a host cell containing said nucleic acid molecule, and a method for screening a test compound for inhibition of MOAT mediated transport by using a host cell comprising SEQ ID No. 5.

Group VIII, claim(s) 30-32, drawn to an antibody specific to MOAT-D.

Group IX, claim(s) 33 and 59, drawn to an oligonucleotide between 10 and 200 nucleotides, which specifically hybridize with a protein translation initiation site in a nucleotide sequence encoding SEQ ID No. 6, and a kit containing oligonucleotide primer sequence for amplification.

Group X, claim(s) 34-40, 45-51 and 56-58, drawn to an isolated nucleic acid molecule having the sequence of SEQ ID No. 7 (MOAT-E), a vector comprising said nucleic acid molecule, a host cell containing said nucleic acid molecule, and a method for screening a test

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compound for inhibition of MOAT mediated transport by using a host cell comprising SEQ ID No. 7.

Group XI, claim(s) 41-43, drawn to an antibody specific to MOAT-E.

Group XII, claim(s) 44 and 59, drawn to an oligonucleotide between 10 and 200 nucleotides, which specifically hybridize with a protein translation initiation site in a nucleotide sequence encoding SEQ ID No. 8, and a kit containing oligonucleotide primer sequence for amplification.

Group XIII, claim(s) 52-55, drawn to a transgenic animal comprising the nucleotide sequence of SEQ ID No. 1 (MOAT-B) or harboring homozygous null mutation of SEQ ID No. 1.

Group XIV, claim(s) 52-55, drawn to a transgenic animal comprising the nucleotide sequence of SEQ ID No. 3 (MOAT-C) or harboring homozygous null mutation of SEQ ID No. 3.

Group XV, claim(s) 52-55, drawn to a transgenic animal comprising the nucleotide sequence of SEQ ID No. 5 (MOAT-D) or harboring homozygous null mutation of SEQ ID No. 5.

Group XVI, claim(s) 52-55, drawn to a transgenic animal comprising the nucleotide sequence of SEQ ID No. 7 (MOAT-E) or harboring homozygous null mutation of SEQ ID No. 7.

2. The inventions listed as Groups I-XVI do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: Groups I, III, VII and X are drawn to different products having different chemical structures, physical properties and biological functions. They

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represent different genes encoding different proteins having different biological functions. Thus, groups I, III, VII and X do not relate to a single general inventive concept under PCT Rule 13.1. Similarly, groups II, IV, VIII and XI are drawn to antibodies that are specific to different proteins, and they do not relate to a single general inventive concept under PCT Rule 13.1. Groups V, VI, IX and XII are drawn to oligonucleotides that specifically hybridize to nucleotide sequences encoding different proteins, and they do not relate to a single general inventive concept under PCT Rule 13.1. Groups XIII-XVI are drawn to transgenic animals having nucleotide sequences encoding different proteins or having null mutation of said nucleotide sequence, and they do not relate to a single general inventive concept under PCT Rule 13.1.

Groups I, III, VII, X, groups II, IV, VIII, XI, groups V, VI, IX, XII, and groups XIII-XVI are drawn to different products: nucleic acids, antibodies, oligonucleotides and transgenic animals. They are different products having different chemical structure and biological functions and different uses. Thus, they do not relate to a single general inventive concept under PCT Rule 13.1.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently

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named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(I).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Shin-Lin Chen whose telephone number is (703) 305-1678. The examiner can normally be reached on Monday to Friday from 9 am to 5:30 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Deborah Reynolds can be reached on (703) 305-4051. The fax phone number for this group is (703) 308-4242.

Any inquiry of a general nature or relating to the status of this application should be directed to the Group receptionist, whose telephone number is (703) 308-0196.

Shin-Lin Chen, Ph.D.

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